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(21) International Application Number: PCT/US99/20547 (22) International Filing Date: 7 September 1999 (07.09.99) (30) Priority Data: 09/160,618 24 September 1998 (24.09.98) US (71) Applicant: DRUG DELIVERY SYSTEMS, INC. [US/US]; 5022 N.W. 104th Way, Coral Springs, FL 33076 (US). (72) Inventor: CHRISTENSEN, Edwin, H.; 5022 N.W. 104th Way, Coral Springs, FL 33076 (US). (74) Agents: SHEKLETON, Gerald, T. et al.; Welsh & Katz, Ltd., 22nd floor, 120 S. Riverside Plaza, Chicago, IL 60606 (US).		(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: SEMI-MOIST ORAL DELIVERY SYSTEM (57) Abstract These and other objects are attained by the subject invention wherein there is provided a carrier or product formed of a matrix having starch, sugar, fat, polyhydric alcohol and water in suitable ratios such that there exists a water activity of 0.6–0.75. The water activity of the product matrix is adjusted up or down so that the availability of water in the finished product is not detrimental to the included active ingredient, be it pharmaceutical, nutraceutical, or a vitamin mineral complex.		

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SEMI-MOIST ORAL DELIVERY SYSTEM**Field of the Invention**

5 This application is directed to a means for delivering pharmaceuticals, nutraceuticals and the like to a mammal and more specifically, the control of the water activity of a food product matrix for use in the incorporation of a pharmaceutical, nutraceutical or other bioactive compound into the matrix.

Background of the Invention

10 Pharmaceutical and nutraceutical products intended for oral administration are typically provided in tablet, capsule, pill, lozenges and caplet form. These products are swallowed whole or chewed in the mouth for delivery of the active ingredient into the alimentary system of a body. Such oral delivery systems are sometimes made chewable to ease drug administration in pediatric and geriatric
15 patients. Such concerns with ease of administration may be amplified when dealing with pets and other animals.

As a result, several approaches have been utilized in formulating oral delivery systems, including gums and candy bases. The use of such delivery systems is limited by the reaction of the active ingredient, whether it be pharmaceutical,
20 nutraceutical or other ingredients, to the existence of water in the system.

Summary of the Invention

Therefore, an object of the subject invention is a method of controlling water activity in an oral delivery system and the product thereof.

A further object of the subject invention is a oral delivery system for
25 pharmaceuticals, nutraceuticals or other active ingredient which matches the water activity of the carrier to the included active ingredient.

Description of the Preferred Embodiments

By the subject invention, a soft chewable oral delivery system is provided. The dosage form may be in tablet form and may contain one or more active ingredients. The active ingredients are incorporated into the system which is described in further detail below and which includes a starch component, a fat or oil, a sugar component, a polyhydric alcohol, water and other minor ingredients. Into this mixture is placed the active ingredient. After mixing and extruding these ingredients, the extrudate is formed into the appropriate shape. The relative proportions of the mixture are as follows.

	Starch	10-50%
	Fat or Oil	0-40%
	Sugar	5-25%
	Polyhydric Alcohol	10-50%
15	Water	5-20%
	Salt (NaCl)	1-5%
	Active Ingredient	0.1-5%

Generally speaking, the starch component of the matrix comprises 10 to 50 percent by weight of the matrix. More particularly, the starch component of the matrix comprises 15 to 40 percent by weight of the matrix.

While starch for use in the matrix can be of any suitable type, it is most preferred that at least part of the starch in the matrix be a highly derivatized or pregelatinized starch. If a highly derivatized starch is present in the matrix, it should be present in an amount of about 1/2 percent by weight of the total starch and the

balance of the starch being non-derivatized. More preferably, about 20-40 percent by weight of the total matrix and about 45% of the total starch should be the derivatized starch. An example of a preferred pregelatinized starch is A.E. Staley's NU-COL 4227 or SOFT-SET.

5 Other amylaceous ingredients may be used in combination with the derivatized starch or alone, provided the starch limits are not exceeded. The amylaceous ingredients can be gelatinized or cooked before or during the forming step to achieve the desired matrix characteristics. If gelatinized starch is used, it may be possible to prepare the product of the subject invention or perform the method of the
10 subject invention without heating or cooking of any sort. However, if ungelatinized (ungelled) or uncooked starch is used, the matrix must be cooked sufficiently to gel or cook the starch to reach the desired content.

 Starches that can serve as a base starch for derivatization include regular corn, waxy corn, potato, tapioca, rice, etc. Such types of derivatizing agents
15 for the starch include but are not limited to ethylene oxide, propylene oxide, acetic anhydride, and succinic anhydride, and other food approved esters or ethers, introducing such chemicals alone or in combination with one another. Prior crosslinking of the starch may or may not be necessary based on the pH of the system and the temperature used to form the product.

20 By "amylaceous ingredients" is meant those food-stuffs containing a preponderance of starch and/or starch-like material. Examples of amylaceous ingredients are cereal grains and meals or flours obtained upon grinding cereal grains such as corn, oats, wheat, milo, barley, rice, and the various milling by-products of these cereal grains such as wheat feed flour, wheat middlings, mixed feed, wheat

shorts, wheat red dog, oat groats, hominy feed, and other such material. Also included as sources of amylaceous ingredients are the tuberous food stuffs such as potatoes, tapioca, and the like.

Another component of the matrix is a fat component such as fat or oil of animal or vegetable origin. Typical animal fats or oils are fish oil, chicken fat, tallow, choice white grease, prime steam lard and mixtures thereof. Other animal fats are also suitable for use in the matrix. Vegetable fats or oils are derived from corn, soy, cottonseed, peanut, flax, rapeseed, sunflower, other oil bearing vegetable seeds, and mixtures thereof. Additionally, a mixture of animal or vegetable oils or fats is suitable for use in the matrix. The fat component of the matrix is about 0 to about 40 % by weight of the matrix. More specifically, the fat component of the matrix is about 20 percent by weight of the matrix.

The polyhydric alcohol component of the matrix can be selected from glycerol, sorbitol, propylene glycol, 1,3-butanediol, and mixtures thereof with each other and other polyhydric alcohols. Generally the polyhydric alcohol comprises about 10 to about 50 percent by weight of the matrix. More specifically, the polyhydric alcohol comprises about 20 to about 40 percent by weight of the matrix.

The sugar component can be employed in a dry or crystalline condition or can be an aqueous syrup having a sugar concentration of from 50 to about 95, preferably from 70 to about 80, weight percent. The sugar used can be lactose, sucrose, fructose, glucose, or maltose, depending on the particular application and price or availability of a particular sugar. Examples of various well established sources of these sugars are, corn syrup solids, malt syrup, hydrolyzed corn starch, hydrol (syrup from glucose manufacturing operations), raw and refined cane and beet

sugars, etc.

Water must be present in the matrix at least about 5 percent by weight of the matrix. More specifically, water is present in the matrix about 5 percent to about 20 percent by weight of the matrix. The matrix thus formed usually has a water activity of 0.60 to 0.75.

While water must be at least 5 percent by weight of the matrix, when the matrix is used in a food product, the moisture of the food product must be adjusted. Generally the moisture content of the matrix is such to give a moisture content of 5-15 percent to the final soft dry food product. More preferred is a moisture content of 5 percent to 14 percent. Most preferred is a moisture content of 8 percent to 13 percent. The desired moisture content may be achieved in any suitable fashion. Normal processing may produce the moisture content desired. A standard drying step is optional and may be used if necessary.

The active ingredient may be any drug, nutrition agent, or the like which can be orally administered. Exemplary of such active ingredients are the following: nutraceuticals, such as chromium picolinate, potassium gluconate and methionine amino acid; prescription drugs, such as ivermectin, fenbendazole, piperazine, magnesium hydroxide, stranzole, furosemide, penicillin, amoxicillin, prednisolone, methylprednisolone, acepromazine; and, other pharmaceutical products, such as aspirin, prozac, zantac, and benedryl. Minor amounts of flavorants, colorants, glycerin, flavor enhancers, sweeteners, emulsifiers, antibitterness agents, taste masking agents, stabilizers, preservatives, or combinations thereof may be added.

To form the matrix, the starch system, fat, polyhydric alcohol, corn

syrup and water are mixed with a screw extruder, permitting addition of ingredients and variable heating at different points along the barrel. Other mixing apparatus, such as a sigma mixer, swept wall heat exchanger or the like may be used. If a coloration is desired in the final product, cooked or pregelled starches are used to form the matrix. The use of these starches avoids high cooking temperatures which would destroy the desired coloration and/or active ingredient. If coloration active temperature sensitivity is not a problem, it is possible to use an uncooked or ungelatinized starch to form the matrix and cook or gel the starch as the process is carried out. The incorporation of a derivatized starch in the product more clearly guarantees the softness of the product for a longer period of time. Softness is also provided by the fats and oils. In this fashion a suitable matrix is provided for use with a wide variety of active ingredients.

Having fully described the invention, the following examples are presented to illustrate the invention without limitation thereof. In these examples all parts percentages are by weight unless otherwise specified.

EXAMPLE 1 -- Carrier

INGREDIENT	PARTS
Regular Corn Starch (Purefood GMI)	18.0
Pregel Starch (SOFT SET)	15.0
Corn Syrup (Star Dri Corn Syrup Solids)	15.0
Corn Oil	20.0
Sorbitol	20.0
H ₂ O	10.0
Salt	2.0
TOTAL	100.0

- The above ingredients are mixed at temperatures of about 125°F, extruded and cut into a suitable tablet size. This product has an oily, bubbly appearance suggesting cutting back on the oil content. Temperature was also adjusted during each of the following examples to eliminate puffing of the product as it exits the extruder.

10 **EXAMPLE 2 -- Guaifenesin**

INGREDIENT	PARTS
Regular Corn Starch (Purefood GMI)	17.9
Pregel Starch (SOFT SET)	15.0
Corn Syrup (Star Dri Corn Syrup Solids)	15.0
Sorbitol	39.3
H ₂ O	10.0
Salt	2.0

Guaifenesin*	0.8
TOTAL	100.0

* Available from Arrow Chemical Co., N.J.

EXAMPLE 3 -- Vitamins

INGREDIENT	PARTS
Regular Corn Starch (Purefood GMI)	17.9
Pregel Starch (SOFT SET)	15.0
Corn Syrup (Star Dri Corn Syrup Solids)	15.0
Sorbitol	35.1
H ₂ O	10.0
Salt	2.0
Vitamin and Mineral Mix*	5.0
TOTAL	100.0

* Commercially available mixture available from Archer Daniels Midland.

5 EXAMPLE 4 -- Flax

INGREDIENT	PARTS
Regular Corn Starch (Purefood GMI)	17.9
Pregel Starch (SOFT SET)	15.0
Corn Syrup (Star Dri Corn Syrup Solids)	15.0
Sorbitol	35.1
H ₂ O	10.0
Salt	2.0
Flax*	5.0
TOTAL	100.0

*Available from Enreco Flax.

EXAMPLE 5 -- Acetaminophen

INGREDIENT	PERCENT
Regular Corn Starch (Purefood GMI)	17.9
Pregel Starch (SOFT SET)	15.0
Corn Syrup (Star Dri Corn Syrup Solids)	15.0
Sorbitol	39.1
H ₂ O	10.0
Salt	2.0
Acetaminophen*	0.8
Red Coloring #40	0.1
Flavoring (Cherry)	0.1
TOTAL	100.0

* Available from Mallinckrodt as Compap

EXAMPLE 6 -- Carrier

INGREDIENT	PARTS
Regular Corn Starch (Purefood GMI)	17.9
Pregel Starch (SOFT SET)	15.0
Corn Syrup (Star Dri Corn Syrup Solids)	15.0
Sorbitol	40.1
H ₂ O	10.0
Salt	2.0
TOTAL	100.0

TABLE 1

Example	Active	Oil/Sugar	A _w	Extrusion Temp.
1	Premix	Corn Oil/Sorbitol	N/A	125
2	Guaifenesin	100% Sorbitol	0.656	115
3	Vitamin Mix	100% Sorbitol	0.651	115
4	Flax	100% Sorbitol	0.673	115
5	Acetaminophen	100% Sorbitol	0.666	115
6	Premix	100% Sorbitol	0.61	115

By the above examples and Table 1 it is apparent that an oral delivery system for the administration for pharmaceuticals, nutraceuticals, vitamins and minerals and other active ingredients may be provided in a chewable form by the subject invention. If the active ingredient is water sensitive such as aspirin, then the amount of polyhydric alcohol is increased, the water activity is depressed to about 0.65 and the stability and texture of the resultant product is maintained. If the active ingredient requires or can tolerate the presence of free water for its activity, such as in the case of Guaifenesin, the amount of polyhydric alcohol may be decreased, while maintaining the level of such polyhydric alcohol such that a soft texture of the resulting tablet is maintained. In the case of Guaifenesin, then an A_w of 0.70 may be utilized and a softer, more chewable texture achieved. An effective oral delivery system in which the texture and stability of the product and activity of the active

ingredient is controllable, is the result.

While the invention has been described with reference to a preferred embodiment, it will be understood by those skilled in the art that various changes may be made and equivalents may be substituted for elements thereof without departing
5 from the scope of the invention. In addition, many modifications may be made to adapt a particular situation or material to the teachings of the invention without departing from the essential scope thereof. Therefore, it is intended that the invention not be limited to the particular embodiment disclosed as the best mode contemplated for carrying out this invention, but that the invention will include all embodiments and
10 equivalents falling within the scope of the appended claims.

Various features of the invention are set forth in the following claims.

What is claimed is:

1. A carrier for the oral administration of an active ingredient to mammals comprising:

10-50% starch,

5 0-40% fat or oil,

8-50% polyhydric alcohol,

5-25% sugar

5-20% water, and

1-5% salt

10 said carrier having an A_w of about 0.60 to 0.75.

2. The carrier of claim 1 wherein the water content is about 10%.

3. The carrier of claim 1 wherein the polyhydric alcohol content is about 40%.

4. The carrier of claim 1 wherein the starch content is about 32%.

5. The carrier of claim 1 wherein the pregelatinized starch content is about 15%.

6. The carrier of claim 1 wherein the active ingredient is a pharmaceutical.

7. The carrier of claim 1 wherein the active ingredient is a nutraceutical.
8. The carrier of claim 1 wherein the active ingredient is a vitamin and mineral mix.
9. The carrier of claim 1 where the A_w is 0.65 and the active ingredient is aspirin.
10. The carrier of claim 1 wherein the polyhydric alcohol is sorbitol.
11. The carrier of claim 1 wherein the sugar content is about 15%.
12. A method of making a carrier for an active ingredient for use in an oral administration of the active ingredient, comprising the steps of:
 - a) forming a matrix by mixing
 - 10-50% starch,
 - 0-40% fat or oil,
 - 10-50% polyhydric alcohol,
 - 5-25% sugar,
 - 5-20% water, and
 - 1-5% salt
 - b) adjusting the relative amounts of polyhydric alcohol and water to control the A_w of said carrier;
whereby the controlled A_w permits the moisture in the carrier to be at a

level not inimical to the active ingredient.

13. The method of claim 12 including adjusting the water content to about 10%.

14. The method of claim 12 including adjusting the polyhydric alcohol content to about 40%.

15. The method of claim 12 including adjusting the starch content to about 32%.

16. The method of claim 12 including adjusting the sugar content to about 15%.

17. The method of claim 12 including adjusting the pregelatinized starch content is about 15%.

18. The method of claim 12 including the step of mixing in a pharmaceutical.

19. The method of claim 12 including the step of mixing in a nutraceutical.

20. The method of claim 12 including the step of mixing in a vitamin and mineral mix.

21. The method of claim 12 including the step of adding sorbitol.

22. The method of claim 12 including the step of controlling the A_w to be at about 0.60 to about 0.75.

23. The method of claim 12 including the step of adding aspirin as the active ingredient and controlling the A_w of said carrier to about 0.65.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US99/20547

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : Please See Extra Sheet.

US CL : Please See Extra Sheet.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : Please See Extra Sheet.

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

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C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4,284,652 A (CHRISTENSEN) 18 August 1981, entire document; col. 7, lines 55-68.	1-8, 10-22
-		-----
Y		10, 23
Y	US 4,025,624 A (ALPHIN et al.) 24 May 1977, col. 8, lines 60-64.	1-23

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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INTERNATIONAL SEARCH REPORT

International application No.

PCT/US99/20547

A. CLASSIFICATION OF SUBJECT MATTER:
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A61K 9/00, 47/00, 9/14

A. CLASSIFICATION OF SUBJECT MATTER:
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B. FIELDS SEARCHED

Minimum documentation searched

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